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(54) **Transportvehicles for macromolecules**

Transportvehikel für Makromoleküle

Véhicules de transport pour macromolécules

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Remarks:

The file contains technical information submitted  
after the application was filed and not included in this  
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**EP 0 755 924 B1**

## Description

[0001] The present invention relates to new compounds which are capable of introducing macromolecules into eucaryotic cells.

[0002] The introduction of macromolecules, including DNA, proteins and the like, into eucaryotic cells can be carried out in different ways, for instance by means of transport vehicles. Such vehicles introduce a molecule into the cell, for instance by means of endocytosis. The vehicles may bind, but for instance also encapsulate, the molecules to be transported. In the latter case the vehicles are referred to as vesicles. Known vesicles are liposomes which consist of a bilayer of phospholipids.

[0003] EP-A516 194 discloses cationic amphiphilic compounds containing pharmaceutical preparations which furthermore contain one monovalent anion and an antiviral compound.

[0004] US-A-4,824,850 describes compounds which are adapted for the site-specific/sustained delivery of centrally acting drug species to the brain.

[0005] Liposomes are for instance used to introduce medicines into the cell. It appeared that liposomes are incorporated into the cell both *in vivo* and *in vitro* by means of endocytosis (Nandi, P.K. et al. (1986) J. Biol. Chem. 261: 16722; Heath, T.D. (1987) Methods Enzymol. 149:111). This means that the largest portion of the material which is incorporated in the cell will ultimately appear in the lysosomal apparatus, where it will be decomposed. Particularly for substances which have their effect in the cytoplasm or the nucleus this is obviously very disadvantageous.

[0006] If the substances to be introduced are hydrophilic it will be difficult to introduce them into liposomes. The main portion of the material remains in the aqueous phase. Particularly in case of expensive substances, like probes and many medicines, this is an obvious disadvantage.

[0007] To prevent that the substances to be introduced into the cell end up in the cell by means of endocytosis, attempts have been made to use fusogenic phospholipids as transport vehicles. The use of fusogenic phospholipids should result in fusion of the from the fusogenic phospholipids formed vesicles with the cell membrane and thus introduce their contents into the cell. However, such attempts have not proven to be very successful because fusogenic liposomes have a strong tendency to mutually merge instead of fusing with the cell membrane (Fontein, T.A.A., Ph. D. Thesis (1992)).

[0008] One of the most important applications in which molecules are introduced into a cell is transfection of the (eucaryotic) cell with DNA or RNA. Transfection is being used for studying the function and regulation of genes and proteins, but also for the genetic modification of microorganisms, plants and animals. There is a large number of artificial techniques which allow DNA to be introduced into a cell, including DNA-micro-injection, DNA-coprecipitation within inorganic salts or with polycations, DNA-encapsulation in liposomes, and making the cell membrane permeable with the aid of chemical or physical means.

[0009] A more recent technique (Felgner, P.L. et al., Proc. Natl. Acad. Sci. U.S.A. 1987 vol. 84, 7413-7417) involves the use of cationic amphiphilic molecules as transport vehicles. One of the best-known amphiphiles is the quaternary ammonium amphiphile DOTMA (N-[1-(2,3-dioleoyloxy)propyl]-N,N,N-trimethylammonium chloride) which in combination with dioleoyl phosphatidyl ethanolamine (DOPE), is commercially available with the name Lipofectine™. Both molecules are lipidic (analogues), which form liposomes, which will form complexes with the negatively charged nucleic acids. Supposedly, the liposomes merge with the plasma membrane and introduce in this way nucleic acids into the cell. However, it could also be done by means of endocytosis. The exact mechanism is yet unknown. With the aid of Lipofectine™ the transfection efficiency may be enhanced by a factor of 30 with respect to other known systems, including the classical calcium phosphate precipitation method. However, the disadvantage of Lipofectine™ is its toxicity and therefore it may be difficult or not possible to use it *in vivo*. Therefore, a demand still remains for other and better transfection methods.

[0010] It is the aim of the present invention to provide new cationic amphiphilic compounds, which allow high efficiencies, for the introduction into a cell of nucleic acids and other macromolecules, including for example proteins and medicines.

[0011] The aim is achieved by the invention by compounds according to claim 1. Preferred compounds are the subject of the claims 2 to 13. A particularly advantageous compound according to the invention is 1-methyl-4-(19-*cis,cis*-heptatriaconta-9,28-dienyl) pyridinium chloride (SAINT-2). The compounds according to the invention are all based on a pyridine ring, which is at one or two positions substituted by a long (ar)alkyl chain. It has been found that with the amphiphiles according to the invention, and particularly with the compound here referred to as "SAINT-2", a transfection efficiency can be obtained which, dependent on the cell type, is at least eight times higher as that of Lipofectine™.

[0012] With SAINT-2/DOPE it also proved to be possible to introduce proteins, particularly gelonine (30kD), into the cell. Other cell types, particularly Baby Hamster Kidney (BHK) cells, may be transfected. This is impossible with Lipofectine™ for BHK cells. SAINT-2/DOPE yields even better results with BHK cells than Lipofectine™ with COS-7 cells.

[0013] The compounds according to the invention may be synthesized in a well-known fashion. The synthesis will be further illustrated in the examples.

[0014] The amphiphiles according to the invention may be used in a large number of applications.

[0015] The transport into the cell of nucleic acids and their derivatives is of importance for transfection. The aim of transfection is, for instance, to make proteins or to perform research. Furthermore, transfected nucleic acids, possibly labelled with streptavidine or radioactively labelled, may be used for in situ hybridisation. A more advanced application is to influence gene expression, for instance blocking of genes by antisense strands. Furthermore, gene expression may also be stimulated. Furthermore, the defect genes may be replaced. The latter two applications are of particular importance in gene therapy.

[0016] The advantage of compounds according to the invention is that they, as compared to the known transport vehicles, can be used in much lower, non-toxic concentrations. Probably, they also do not cause an immunologic response.

[0017] If DNA and/or RNA are to be introduced into a cell the compounds and the nucleic acids have to be mixed in a certain ratio. It has been found that for the known amphiphiles, including DOTMA, there exists an optimum amphiphile concentration (Felgner, P.L. et al. (1987) Proc. Natl. Acad. Sci. USA 84:7413). The transfection efficiency again reduces if a certain amount is exceeded. A comparable situation also holds for the compounds according to the invention.

[0018] The cationic amphiphiles according to the invention may also be used to transport negatively charged proteins, including gelonine in particular, into the cell.

[0019] The amphiphiles may also be used to transport substances like cytostatics. Lipophilic cytostatics in particular do interact with the compounds according to the invention and may in this way be introduced into the cell very efficiently.

[0020] In a preferred embodiment of the invention the transport vehicles may be purposely brought to a specific site by mixing the amphiphiles with a targeting molecule, such as, for instance, an antibody which is directed against an epitope in the neighbourhood of the site where the incorporated substance has to exercise its activity. The antibody is preferably coupled to the amphiphilic compound but it may also be coupled, for instance, through a spacer, to the substance to be transported. In order to facilitate the translocations of DNA or other macromolecules across the cell membrane the compounds according to the invention may also be mixed with a phospholipid or with each other.

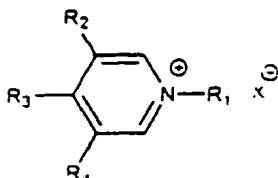
[0021] The present invention will be illustrated in further detail with by means of the accompanying examples which are only serve as an illustration and do not limit the scope of the invention.

## EXAMPLES

### EXAMPLE 1

#### Synthesis

[0022] Compounds with the general structure formula



may be divided in a number of groups dependent on their substituents. The synthesis of four of those groups will be given below as an example.

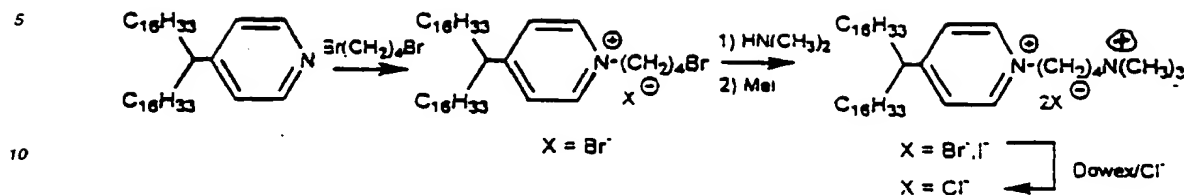
#### 1. 4-Substituted N-alkylpyridinium salts.

##### 1.1. Synthesis of 1-methyl-4-(1-octadecylnonadecyl)pyridinium chloride.

[0023] The compound is synthesised according to scheme 1 below as described by E.J.R. Sudhölter in his Ph.D. thesis at the University of Groningen, 1981, page 37.



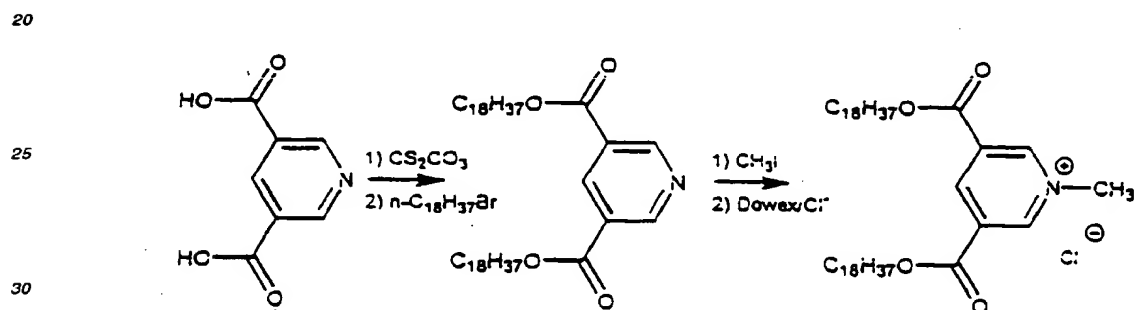
Scheme 3:



## 2. 3,5-Disubstituted-N-alkylpyridinium salts.

15 **[0032]** The general synthesis according to scheme 4 below was described in the literature by Sudhölter (*vide supra*) and Wang et al., J. Org. Chem. 42, 1286 (1977).

Scheme 4:



## 2.1. 1-methyl-3,5-dicarbo-N-octadecyloxy pyridinium chloride.

35 **[0033]** This compound was synthesized according to scheme 4.

**[0034]** NMR data: <sup>1</sup>H NMR(CDCl<sub>3</sub>): δ 0.85 (t, 6H); 1.30 (chain, 64H); 4.40 (t, 4H); 5.03 (s, 3H); 9.20 (t, 1H); 10.00 (d, 2H).

## 3. 4-Substituted-N-alkyl pyridinium salts.

**[0035]** The synthesis was described by F.J.A. Hundscheid and J.B.F.N. Engberts, J. Org. Chem. 49, 3088 (1984).

## 3.1. 1-Methyl-4((-n-hexadecyloxy)carbonyl) pyridinium iodide.

45 **[0036]** The synthesis of this compound and its characterisation are described by Hundscheid and Engberts (*vide supra*).

**[0037]** NMR data: <sup>1</sup>H NMR(CDCl<sub>3</sub>): δ 0.9 (t, 3H); 1.25 (m, 28H); 4.35 (t, 2H); 4.70 (s, 3H); 8.35 (d, 2H); 9.35 (d, 2H).

## 4. 4-Substituted-N-aralkyl pyridinium salts.

## 4.1. 1-(3-phenyl-1-propyl)-4-n-dodecylpyridinium iodide.

55 **[0038]** The compound was synthesized by boiling a mixture of 2.26 g (9.2 mmol) 1-iodo-3-phenyl propane and 2.57 g (10.0 mmol) 4-n-dodecylpyridine in 35 ml of dry acetone for 16 hours. The solvent was evaporated and the yellow solid substance was recrystallized from THF/ether. The yield is 3.06 g (6.2 mmol), melting point 79.0-80.0°C.

**[0039]** NMR data: <sup>1</sup>H NMR(CDCl<sub>3</sub>): δ 0.83 (t, 3H); 1.21 (chain, 20H); 1.61 (m, 2H); 2.36 (m, 2H); 2.78 (m, 2H); 4.89 (t, 2H); 7.05-7.20 (m, 5H); 7.73 (d, 2H); 9.30 (d, 2H).

## EXAMPLE 2

Formation of unilamellar vesicles

[0040] A suitable amount of lipid was dried under N<sub>2</sub>(g). In case of combinations of substances these are first mixed and then dried. The lipid film layer is subsequently dried further under vacuum. The lipids are then suspended, vortexed and subsequently sonicated in a suitable volume of water until the solution is clear.

## EXAMPLE 3

Transfection of eucaryotic cells by compounds according to the invention

[0041] DNA and unilamellar vesicles, as prepared in Example 2, are both brought into Hepes buffered saline (HBS, pH 7.4; both 0.5 ml) and subsequently mixed. The DNA/amphiphile complex is directly formed. In a typical transfection experiment 1 µg of DNA and 7.5-10 µg of the amphiphile SAINT-2 (1-methyl-4-(19-cis-cis-heptatriacontadienyl-9-28)pyridinium chloride) or 1 µg of DNA and 10-15 µg of total amphiphile (SAINT-2/DOPE 1:1) is used.

[0042] Cells in six-well plates, which are confluent by 70-80%, are washed twice with 1 ml of HBS and subsequently 1 ml of the DNA/amphiphile complex was added per well. The cells were incubated during 4 hours at 37°C after which 1 ml Dulbecco Modified Eagle Medium (DMEM) supplemented with 10% Foetal Calf Serum (FCS) was added. After an incubation of 16 hours at 37°C the medium was exchanged by 2 ml fresh DMEM with 10% FCS. After a subsequent incubation of 28 hours at 37°C the cells were gathered. The cells were washed twice with a phosphate buffered saline (PBS) and scraped in 300 µl 1x lysis buffer (Promega). The scraped cells were incubated for 10 minutes at 56°C and subsequently centrifuged at maximum speed for two minutes at room temperature. On the supernatant an enzyme determination (CAT-assay) and a protein determination (Lowry) were carried out.

[0043] 100 µl of the cell extract was incubated together with 3 µl <sup>14</sup>C-chloramphenicol (25 mCi/l), 5 µl N-butyryl-CoA (2 mg/ml) and 17 µl 0.25 M Tris.HCL (pH 8.0) during 90 minutes at 37°C. The reaction was stopped by adding 0.3 ml of mixed xylenes (Aldrich). The samples were vortexed for 30 seconds and subsequently centrifuged at maximum speed for 3 minutes at room temperature. The organic phase was again extracted with 0.1 ml 0.25 M Tris.HCL, vortexed for 30 seconds and centrifuged for 3 minutes. 4 ml of counting fluid was added to 0.2 ml of the organic phase and the radio activity was measured.

[0044] It was found that transfection of COS-7 cells with the new amphiphile (SAINT-2 and SAINT-2/DOPE) is eight times more efficient than that with DOTMA/DOPE vesicles (see Fig. 1).

[0045] It appeared that with the new amphiphile also other cell types, including for instance BHK cells, can be transfected (Fig. 2). When a stable transfection is carried out with the new amphiphile it appeared to be possible to transfect 42-45% of the COS-7 cells. With DOTMA-DOPE vesicles on average 25-29% of the cells are transfected.

## EXAMPLE 4

Transport of proteins in an eucaryotic cell

[0046] The synthetic amphiphile SAINT-2 is, in combination with DOPE, a suitable agent for the delivery of proteins into cells. The efficiency of protein internalisation with SAINT-2/DOPE as a carrier can be monitored with the aid of the gelonine protein. Internalized gelonine specifically inhibits the protein synthesis of cells and this inhibition is a direct measure for the amount of gelonine which has been brought into the cell. Unilamellar vesicles of the synthetic agent amphiphile SAINT-2 and DOPE are obtained by bath sonication. Gelonine is added to a certain concentration (0-20 µM) SAINT-2/DOPE in HBS from a stock solution (2 mg/ml).

[0047] CV-1 cells, grown in twelve-well plates, are washed three times with HBS. Subsequently, the cells are incubated for 1 hour at 37°C with the amphiphile/gelonine complex in HBS obtained in this way. After this the cells are again washed three times with HBS.

[0048] The inhibition of protein synthesis by gelonine is being followed by determining the building-in of radioactively labelled methionine into the treated cells. This is carried out by incubating the cells for 30 minutes with 1 µCi <sup>35</sup>S-methionine. Subsequently, the cells are washed three times with PBS and finally scraped in 10% TCA. The cell lysate obtained in this way is washed three times with 10% TCA and the amount of radioactive methionine present in the cell lysate is determined with the aid of a scintillation counter.

[0049] Incubation of CV-1 cells with the amphiphile/gelonine complex gives a strong inhibition of the protein synthesis with respect to the control experiment in which the cells were incubated with the synthetic amphiphile only. At a concentration of 5 µM SAINT-2/DOPE and 1.6 µM gelonine an inhibition of protein synthesis of 50% was obtained.

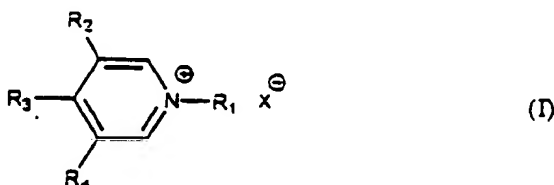
## EXAMPLE 5

Toxicity studies

- 5 [0050] To determine the toxicity of the compound SAINT-2 according to the invention with respect to DOTMA-DOPE the COS-7 cells are incubated with different concentrations of both lipid samples. The residual protein content is taken as a measure for the amount of surviving cells.
- [0051] A decrease of the protein content from 2 to 1 mg/ml was observed for DOTMA-DOPE starting from 71  $\mu$ M lipid. For SAINT-2 a decrease from 2 to 1.75 mg/ml was found starting from 90  $\mu$ M.
- 10 [0052] This shows that SAINT-2 is clearly less toxic than DOTMA-DOPE.

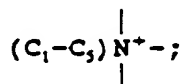
## Claims

- 15 1. Compounds with the general formula I



in which:

$R_1$  is a (C<sub>1</sub>-C<sub>5</sub>)alkyl, ar(alkyl) or an alkyl group with a cationic functional group, like

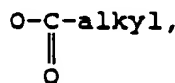


or

$R_1$  is (C<sub>1</sub>-C<sub>5</sub> alkylene) $R_5$  in which  $R_5$  is a structure with the general formula I except  $R_1$

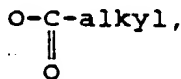
X is a halide counter ion; chosen from Cl<sup>-</sup>, I<sup>-</sup>, Br<sup>-</sup>; and in which

or  $R_3$  is hydrogen and  $R_2$  and  $R_4$  are identical or different and are chosen from the group, comprising branched or linear (C<sub>10</sub>-C<sub>20</sub>)alkyl, a mono- or polyunsaturated (C<sub>10</sub>-C<sub>20</sub>) alkenyl, O=C-O-alkyl,



or ar(alkyl),

or  $R_2$  and  $R_4$  are hydrogen and  $R_3$  is  $-\text{CH}(R_5)_2$  with  $R_5$ , comprising (C<sub>10</sub>-C<sub>20</sub>)alkyl, mono- or polyunsaturated (C<sub>10</sub>-C<sub>20</sub>) alkenyl, O=C-O-alkyl,



or aralkyl,

wherein disclaimed are the compounds with the general formula I in which  $R_1$  is CH<sub>3</sub>,  $R_2$  and  $R_4$  are hydrogen,  $R_3$  is (C<sub>16</sub>H<sub>33</sub>)<sub>2</sub>CH and X is all mentioned counter ions (Cl<sup>-</sup>, I<sup>-</sup>, Br<sup>-</sup>) and disclaimed are the compounds in which  $R_1$  is CH<sub>3</sub>,  $R_2$  and  $R_4$  are C<sub>16</sub>H<sub>33</sub>-O-C(O),  $R_3$  is hydrogen and X is all mentioned counter ions (Cl<sup>-</sup>, I<sup>-</sup>, Br<sup>-</sup>), and

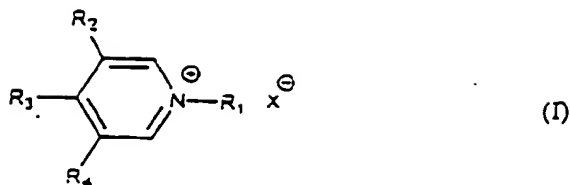
3,5-bis[(decyloxy)-carbonyl]-1-methyl-pyridinium iodide.

2. Compounds according to claim 1, **characterized** in, that  $R_1$  is  $\text{CH}_3$ ,  $R_2$  and  $R_4$  are hydrogen,  $R_3$  is  $(\text{C}_{18}\text{H}_{37})_2\text{CH}$  and X is Cl<sup>-</sup>, Br<sup>-</sup>, I<sup>-</sup>.
3. Compounds according to claim 1, **characterized** in that  $R_1$  is  $\text{CH}_3$ ,  $R_2$  and  $R_4$  are hydrogen,  $R_3$  is  $(\text{C}_{18}\text{H}_{35})_2\text{CH}$  and X is Cl<sup>-</sup>, Br<sup>-</sup>, I<sup>-</sup>.
4. Compounds according to claim 1, **characterized** in that  $R_1$  is  $(\text{CH}_2)_4\text{-N}^+(\text{CH}_3)_3$ ,  $R_2$  and  $R_4$  are hydrogen,  $R_3$  is  $(\text{C}_{18}\text{H}_{35})_2\text{CH}$  and X is Cl<sup>-</sup>, Br<sup>-</sup>, I<sup>-</sup>.
5. Compounds according to claim 1, **characterized** in that  $R_1$  is  $\text{CH}_3$ ,  $R_2$  and  $R_4$  are  $\text{C}_{18}\text{H}_{37}\text{-O-C(O)}$ ,  $R_3$  is hydrogen and X is Cl<sup>-</sup>, Br<sup>-</sup>, I<sup>-</sup>.
6. Compounds according to claim 1, **characterized** in that  $R_1$  is  $\text{CH}_3$ ,  $R_2$  and  $R_4$  are  $\text{R}_5\text{-O-C(O)}$ , in which  $\text{R}_5$  is a saturated  $\text{C}_{10}\text{-C}_{20}$  aliphatic chain,  $R_3$  is hydrogen and X is Cl<sup>-</sup>, Br<sup>-</sup>, I<sup>-</sup>.
7. Compounds according to claim 1, **characterized** in that  $R_1$  is  $\text{CH}_3$ ,  $R_2$  and  $R_4$  are not identical and each represents a group with the formula  $\text{R}_5\text{-O-C(O)}$ , in which  $\text{R}_5$  is a  $\text{C}_{10}\text{-C}_{20}$  alkyl, is,  $R_3$  is hydrogen and X is Cl<sup>-</sup>, Br<sup>-</sup>, I<sup>-</sup>.
8. Compounds according to claim 1, **characterized** in that  $R_1$  is  $\text{CH}_3$ ,  $R_3$  is a group with the formula  $\text{R}_5\text{-O-C(O)}$ , in which  $\text{R}_5$  is a  $\text{C}_{10}\text{-C}_{20}$  alkyl,  $R_2$  and  $R_4$  are hydrogen and X is Cl<sup>-</sup>, Br<sup>-</sup> or I<sup>-</sup>.
9. Compounds according to claim 8, **characterized** in that  $\text{R}_5$  is  $\text{C}_{16}\text{H}_{33}$ .
10. Compounds according to claim 1, **characterized** in that  $R_1$  is  $(\text{CH}_2)_n\text{-C}_6\text{H}_5$ , in which  $n = 3\text{-}6$ ,  $R_2$  and  $R_4$  are hydrogen,  $R_3$  is alkyl or alkenyl and X is Cl<sup>-</sup>, Br<sup>-</sup> or I<sup>-</sup>.
11. Compound according to claim 10, **characterized** in that  $R_1$  is  $(\text{CH}_2)_3\text{-C}_6\text{H}_5$ ,  $R_2$  and  $R_4$  are hydrogen,  $R_3$  is  $n\text{-C}_{12}\text{H}_{25}$  and X is Cl<sup>-</sup>, Br<sup>-</sup> or I<sup>-</sup>.
12. Compounds according to claim 1, **characterized** in that  $R_1$  is  $(\text{CH}_2)_4$ ,  $R_2$  and  $R_4$  are hydrogen,  $R_3$  is  $(\text{C}_{18}\text{H}_{35})_2\text{CH}$ , X is Cl<sup>-</sup>, Br<sup>-</sup>, I<sup>-</sup> and  $\text{R}_5$  is the group with the general formula I, bound through  $\text{R}_1$ , in which  $\text{R}_2$  and  $R_4$  are hydrogen,  $\text{R}_3$  is  $(\text{C}_{18}\text{H}_{35})_2\text{CH}$  and X is Cl<sup>-</sup>, Br<sup>-</sup>, I<sup>-</sup>.
13. Compounds according to one of the preceding claims for use as tool to introduce macromolecules into cells.
14. Composition to introduce macromolecules into cells, comprising vesicles or another type of aggregate formed by at least one compound according to one of the preceding claims in a solvent.
15. Composition according to claim 14, **characterized** in that at least one macromolecule is incorporated in and/or attached to the vesicles.
16. Composition according to claim 14 or 15, **characterized** in that furthermore at least one targeting molecule is attached to the vesicles.
17. Composition according to claim 15, **characterized** in that the targeting molecule is an antibody.
18. Composition according to claim 16, **characterized** in that the antibody is radioactively labelled or labelled with streptavidine.

#### Patentansprüche

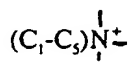
1. Verbindungen mit der allgemeinen Formel I





10 in der:

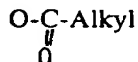
$R_1$  eine  $(C_1-C_5)$ Alkyl-, Ar(alkyl)- oder eine Alkyl-Gruppe mit einer kationischen funktionellen Gruppe wie



ist; oder in der  $R_1$   $(C_1-C_5)$ -Alkylen)  $R_5$  ist, worin  $R_5$  eine Struktur mit der allgemeinen Formel I ohne  $R_1$  ist,

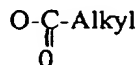
$X$  ein Halogenid-Gegenion ist, das ausgewählt ist aus  $Cl^-$ ,  $I^-$ ,  $Br^-$ ; und in der

20  $R_3$  Wasserstoff ist und  $R_2$  und  $R_4$  gleich oder verschieden sind und ausgewählt sind aus der Gruppe, die verzweigtes oder lineares  $(C_{10}-C_{20})$ Alkyl, ein mono- oder polyungesättigtes  $(C_{10}-C_{20})$ Alkenyl,  $O=C-O$ -Alkyl,



oder Ar(alkyl) aufweist,

oder in der  $R_2$  und  $R_4$  Wasserstoff sind und  $R_3-CH(R_5)_2$  ist, wobei  $R_5$   $(C_{10}-C_{20})$ Alkyl, mono- oder polyungesättigtes  $(C_{10}-C_{20})$ Alkenyl,  $O=C-O$ -Alkyl,



oder Aralkyl aufweist,

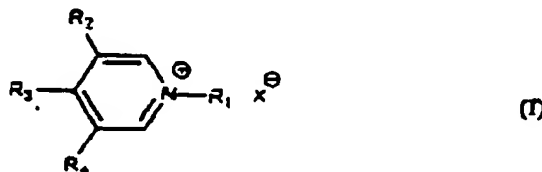
35 unter Ausschluß der Verbindungen mit der allgemeinen Formel I, in denen  $R_1$   $CH_3$  ist,  $R_2$  und  $R_4$  Wasserstoff sind,  $R_3$   $(C_{16}H_{33})_2CH$  ist und  $X$  alle aufgeführten Gegenionen ( $Cl^-$ ,  $I^-$ ,  $Br^-$ ) ist, und unter Ausschluß der Verbindungen, in denen  $R_1$   $CH_3$  ist,  $R_2$  und  $R_4$   $C_{16}H_{33}-O-C(O)$  sind,  $R_3$  Wasserstoff ist und  $X$  alle aufgeführten Gegenionen ( $Cl^-$ ,  $I^-$ ,  $Br^-$ ) ist, und 3,5-Bis[(decyloxy)-carbonyl]-1-methyl-pyridinium-iodid.

- 40
2. Verbindungen nach Anspruch 1, **dadurch gekennzeichnet**, daß  $R_1$   $CH_3$  ist,  $R_2$  und  $R_4$  Wasserstoff sind,  $R_3$   $(C_{16}H_{37})_2CH$  ist und  $X$   $Cl^-$ ,  $Br^-$ ,  $I^-$  ist.
  3. Verbindungen nach Anspruch 1, **dadurch gekennzeichnet**, daß  $R_1$   $CH_3$  ist,  $R_2$  und  $R_4$  Wasserstoff sind,  $R_3$   $(C_{16}H_{35})_2CH$  ist und  $X$   $Cl^-$ ,  $Br^-$ ,  $I^-$  ist.
  4. Verbindungen nach Anspruch 1, **dadurch gekennzeichnet**, daß  $R_1$   $(CH_3)_4-N^+(CH_3)_3$  ist,  $R_2$  und  $R_4$  Wasserstoff sind,  $R_3$   $(C_{18}H_{35})_2CH$  ist und  $X$   $Cl^-$ ,  $Br^-$ ,  $I^-$  ist.
  - 50 5. Verbindungen nach Anspruch 1, **dadurch gekennzeichnet**, daß  $R_1$   $CH_3$  ist,  $R_2$  und  $R_4$   $C_{18}H_{37}-O-C(O)$  sind,  $R_3$  Wasserstoff ist und  $X$   $Cl^-$ ,  $Br^-$ ,  $I^-$  ist.
  6. Verbindungen nach Anspruch 1, **dadurch gekennzeichnet**, daß  $R_1$   $CH_3$  ist,  $R_2$  und  $R_4$   $R_5-O-C(O)$  sind, worin  $R_5$  eine gesättigte, aliphatische  $C_{10}-C_{20}$ -Kette ist,  $R_3$  Wasserstoff ist und  $X$   $Cl^-$ ,  $Br^-$ ,  $I^-$  ist.
  - 55 7. Verbindungen nach Anspruch 1, **dadurch gekennzeichnet**, daß  $R_1$   $CH_3$  ist,  $R_2$  und  $R_4$  nicht gleich sind und jedes eine Gruppe mit der Formel  $R_5-O-C(O)$  darstellt, worin  $R_5$  ein  $C_{10}-C_{20}$ -Alkyl ist,  $R_3$  Wasserstoff ist und  $X$   $Cl^-$ ,  $Br^-$ ,  $I^-$  ist.

8. Verbindungen nach Anspruch 1, **dadurch gekennzeichnet**, daß  $R_1$   $\text{CH}_3$  ist,  $R_3$  eine Gruppe mit der Formel  $R_5\text{-O-C(O)}$  ist, worin  $R_5$  ein  $\text{C}_{10}\text{-C}_{20}$ -Alkyl ist,  $R_2$  und  $R_4$  Wasserstoff sind und  $X$   $\text{Cl}^-$ ,  $\text{Br}^-$  oder  $\text{I}^-$  ist.
9. Verbindungen nach Anspruch 8, **dadurch gekennzeichnet**, daß  $R_5$   $\text{C}_{16}\text{H}_{33}$  ist.
10. Verbindungen nach Anspruch 1, **dadurch gekennzeichnet**, daß  $R_1$   $(\text{CH}_2)_n\text{-C}_6\text{H}_5$  ist, worin  $n = 3\text{-}6$ ,  $R_2$  und  $R_4$  Wasserstoff sind,  $R_3$  Alkyl oder Alkenyl ist und  $X$   $\text{Cl}^-$ ,  $\text{Br}^-$  oder  $\text{I}^-$  ist.
11. Verbindung nach Anspruch 10, **dadurch gekennzeichnet**, daß  $R_1$   $(\text{CH}_2)_3\text{-C}_6\text{H}_5$  ist,  $R_2$  und  $R_4$  Wasserstoff sind,  $R_3$   $n\text{-C}_{12}\text{H}_{25}$  ist und  $X$   $\text{Cl}^-$ ,  $\text{Br}^-$  oder  $\text{I}^-$  ist.
12. Verbindungen nach Anspruch 1, **dadurch gekennzeichnet**, daß  $R_1$   $(\text{CH}_2)_4$  ist,  $R_2$  und  $R_4$  Wasserstoff sind,  $R_3$   $(\text{C}_{18}\text{H}_{35})_2\text{CH}$  ist,  $X$   $\text{Cl}^-$ ,  $\text{Br}^-$ ,  $\text{I}^-$  ist, und  $R_5$  die Gruppe mit der allgemeinen Formel I ist, die über  $R_1$  gebunden ist, worin  $R_2$  und  $R_4$  Wasserstoff sind,  $R_3$   $(\text{C}_{18}\text{H}_{35})_2\text{CH}$  ist und  $X$   $\text{Cl}^-$ ,  $\text{Br}^-$ ,  $\text{I}^-$  ist.
13. Verbindungen nach einem der vorangehenden Ansprüche zur Verwendung als ein Hilfsmittel zum Einführen von Makromolekülen in Zellen.
14. Zusammensetzung zur Einführung von Makromolekülen in Zellen, die Vesikel oder eine andere Art von mittels mindestens einer Verbindung gemäß einem der vorangehenden Ansprüche gebildetem Aggregat in einem Lösungsmittel aufweist.
15. Zusammensetzung nach Anspruch 14, **dadurch gekennzeichnet**, daß mindestens ein Makromolekül in die Vesikel inkorporiert und/oder an die Vesikel gebunden ist.
16. Zusammensetzung nach Anspruch 14 oder 15, **dadurch gekennzeichnet**, daß außerdem mindestens ein Zielfindungsmolekül an die Vesikel gebunden ist.
17. Zusammensetzung nach Anspruch 15, **dadurch gekennzeichnet**, daß Zielfindungsmolekül ein Antikörper ist.
18. Zusammensetzung nach Anspruch 16, **dadurch gekennzeichnet**, daß der Antikörper radioaktiv markiert oder mit Streptavidin markiert ist.

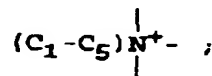
## Revendications

1. Composés répondant à la formule générale I



dans laquelle :

$R_1$  représente un groupe alkyle en  $\text{C}_1$  à  $\text{C}_5$ , Ar(alkyle) ou un groupe alkyle avec un groupe fonctionnel cationique, tel qu'un groupe

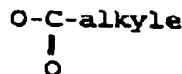


ou bien

$R_1$  représente un groupe (alkylène en  $\text{C}_1$  à  $\text{C}_5$ ) $R_5$  dans lequel  $R_5$  représente une structure de la formule

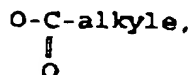
générale I, à l'exception de  $R_1$  ;

X représente un ion complémentaire halogénure, choisi entre  $Cl^-$ ,  $I^-$  et  $Br^-$  ; et dans laquelle soit  $R_3$  représente l'hydrogène et  $R_2$  et  $R_4$  sont identiques ou différents et sont choisis dans le groupe comprenant des radicaux alkyle en  $C_{10}$  à  $C_{20}$  ramifiés ou linéaires, un radical alcényle en  $C_{10}$  à  $C_{20}$  mono- ou polyinsaturé, un radical  $O=C-O$ -alkyle, un radical



ou un radical  $Ar(alkyle)$ ,

soit  $R_2$  et  $R_4$  représentent l'hydrogène et  $R_3$  représente un groupe  $-CH(R_5)_2$  dans lequel  $R_5$  comprend un groupe alkyle en  $C_{10}$  à  $C_{20}$ , alcényle en  $C_{10}$  à  $C_{20}$  mono- ou polyinsaturé,  $O=C-O$ -alkyle,



ou aralkyle,

à l'exception des composés de formule générale I dans laquelle  $R_1$  représente un groupe  $CH_3$ ,  $R_2$  et  $R_4$  représentent l'hydrogène,  $R_3$  représente un groupe  $(C_{16}H_{33})_2CH$  et X représente tous les ions complémentaires précités ( $Cl^-$ ,  $I^-$  et  $Br^-$ ), et

à l'exception des composés dans lesquels  $R_1$  représente un groupe  $CH_3$ ,  $R_2$  et  $R_4$  représentent des groupes  $C_{16}H_{33}-O-C(O)$ ,  $R_3$  représente l'hydrogène et X représente tous les ions complémentaires précités ( $Cl^-$ ,  $I^-$  et  $Br^-$ ) et de l'iodure de 3,5-bis [(décyloxy)-carbonyl]-1-méthyl-pyridinium.

2. Composés suivant la revendication 1, caractérisés en ce que  $R_1$  représente un groupe  $CH_3$ ,  $R_2$  et  $R_4$  représentent l'hydrogène,  $R_3$  représente un groupe  $(C_{16}H_{37})_2CH$  et X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$ .
3. Composés suivant la revendication 1, caractérisés en ce que  $R_1$  représente un groupe  $CH_3$ ,  $R_2$  et  $R_4$  représentent l'hydrogène,  $R_3$  représente un groupe  $(C_{16}H_{35})_2CH$  et X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$ .
4. Composés suivant la revendication 1, caractérisés en ce que  $R_1$  représente un groupe  $(CH_2)_4-N^+(CH_3)_3$ ,  $R_2$  et  $R_4$  représentent l'hydrogène,  $R_3$  représente un groupe  $(C_{16}H_{35})_2CH$  et X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$ .
5. Composés suivant la revendication 1, caractérisés en ce que  $R_1$  représente un groupe  $CH_3$ ,  $R_2$  et  $R_4$  représentent un groupe  $(C_{18}H_{37}-O-C(O))$ ,  $R_3$  représente l'hydrogène et X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$ .
6. Composés suivant la revendication 1, caractérisés en ce que  $R_1$  représente un groupe  $CH_3$ ,  $R_2$  et  $R_4$  représentent des groupes  $R_5-O-C(O)$  dans lesquels  $R_5$  représente une chaîne aliphatique en  $C_{10}$  à  $C_{20}$  saturée,  $R_3$  représente l'hydrogène et X représente un groupe  $Cl^-$ ,  $Br^-$  ou  $I^-$ .
7. Composés suivant la revendication 1, caractérisés en ce que  $R_1$  représente un groupe  $CH_3$ ,  $R_2$  et  $R_4$  ne sont pas identiques et représentent chacun un groupe de formule  $R_5-O-C(O)$ , dans laquelle  $R_5$  représente un groupe alkyle en  $C_{10}$  à  $C_{20}$ ,  $R_3$  représente l'hydrogène et X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$ .
8. Composés suivant la revendication 1, caractérisés en ce que  $R_1$  représente un groupe  $CH_3$ ,  $R_3$  représente un groupe de formule  $R_5-O-C(O)$ , dans laquelle  $R_5$  représente un groupe alkyle en  $C_{10}$  à  $C_{20}$ ,  $R_2$  et  $R_4$  représentent l'hydrogène et X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$ .
9. Composés suivant la revendication 8, caractérisés en ce que  $R_5$  représente un groupe  $C_{16}H_{33}$ .
10. Composés suivant la revendication 1, caractérisés en ce que  $R_1$  représente un groupe  $(CH_2)_n-C_6H_5$  dans lequel n a une valeur de 3 à 6,  $R_2$  et  $R_4$  représentent l'hydrogène,  $R_3$  représente un groupe alkyle ou alcényle et X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$ .

11. Composés suivant la revendication 10, caractérisés en ce que  $R_1$  représente un groupe  $(CH_2)_3-C_6H_5$ ,  $R_2$  et  $R_4$  représentent l'hydrogène,  $R_3$  représente un groupe  $n-C_{12}H_{25}$  et X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$ .
- 5 12. Composés suivant la revendication 11, caractérisés en ce que  $R_1$  représente un groupe  $(CH_2)_4$ ,  $R_2$  et  $R_4$  représentent l'hydrogène,  $R_3$  représente un groupe  $(C_{18}H_{35})_2CH$ , X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$  et  $R_5$  représente le groupe de la formule générale I, lié par  $R_1$ , dans lequel  $R_2$  et  $R_4$  représentent l'hydrogène,  $R_3$  représente un groupe  $(C_{18}H_{35})_2CH$ , et X représente  $Cl^-$ ,  $Br^-$  ou  $I^-$ .
- 10 13. Composés suivant l'une des revendications précédentes, destinés à être utilisés comme outils pour introduire des macromolécules dans les cellules.
14. Composition pour introduire des macromolécules dans des cellules, comprenant des vésicules ou un autre type d'agrégat, formés par au moins un composé suivant l'une des revendications précédentes dans un solvant.
- 15 15. Composition suivant la revendication 14, caractérisée en ce qu'au moins une macromolécule est incorporée et/ou fixée aux vésicules.
16. Composition suivant la revendication 14 ou 15, caractérisée en ce que, en outre, au moins une molécule d'acheminement vers une cible est fixée aux vésicules.
- 20 17. Composition suivant la revendication 15, caractérisée en ce que la molécule d'acheminement vers une cible est un anticorps.
- 25 18. Composition suivant la revendication 16, caractérisée en ce que l'anticorps est marquée avec un marqueur radioactif ou est marquée avec de la streptavidine.

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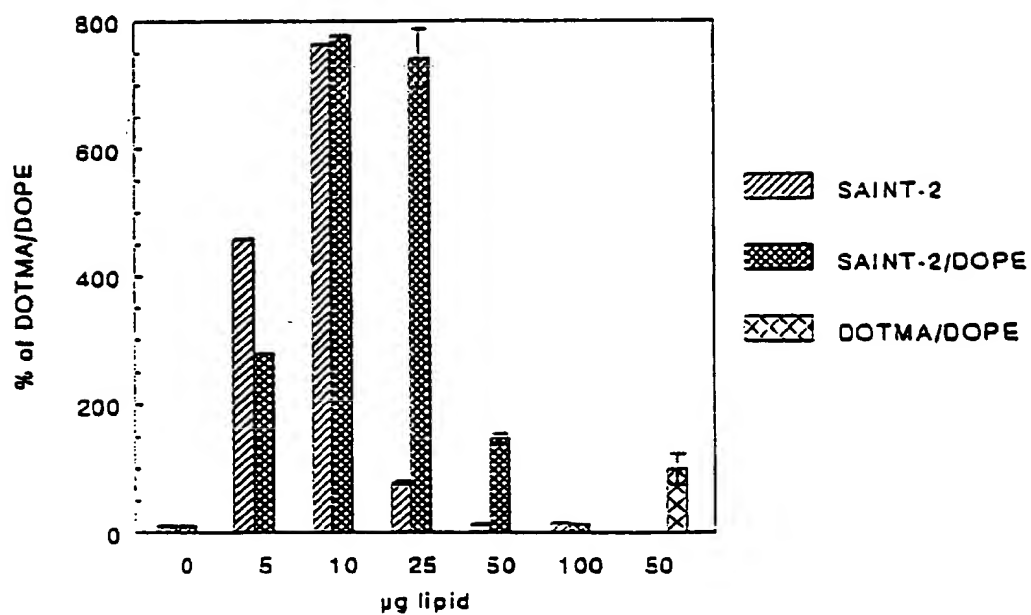
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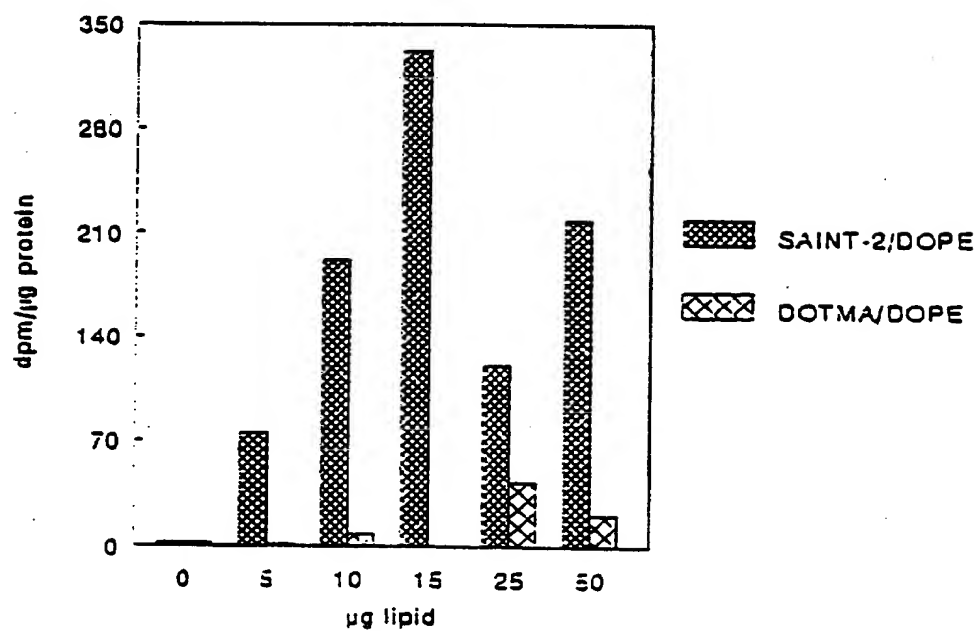
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Transfection of COS-7 cells using  
different synthetic amphiphiles



Transfection of BHK cells using  
different synthetic amphiphiles



# Vesicle-mediated versus CaP-mediated transfection

